



**Bando 2019 - Programma 5 per mille anno 2019  
Investigator Grant (IG)**

**TRANSLATIONAL RESEARCH**

LILT will support research projects in the field of cancer aimed at improving cancer diagnosis and treatment. Particularly considered will be those translational research projects that promise short-medium term effects in clinical practice, concerning new diagnostic methodologies and new therapies. Multicentric studies with national coordination, aimed at validating new diagnostic methods, diagnostic, prognostic and predictive tumor markers, able to improve the clinical management of cancer patients are potentially eligible for funding. Specific research projects on new oncological therapeutic approaches are also eligible for LILT funding as IG. For this type of grants it is necessary to demonstrate solid preliminary experimental data supported by a rigorous biological rationale.

**1. Principal investigator's full name and qualification:**

(Please include: CV in European format with list of publications; IF end Hi-index )

2. Proposal title.....

3. Primary area of Relevance.....

4. Relevance for the National Health System.....

5. Institution / University..... address.....phone..... e-mail.....

6. Authorized Administrative Official.....address.....phone..... e-mail.....

7. Proponent's signature.....

8. Authorized Administrative Official's signature.....

9. Place and date...../.../....

## SELF EVALUATION FORM

1. Investigator's full name: (PI) .....
2. Total papers..... IF.....
3. Total papers (last 10 years).....IF.....
4. Total Papers as first/last author or corresponding author.....
5. Total H-index .....

## PROPOSAL MAIN BODY

<ol style="list-style-type: none"> <li>1. Proposal title.....</li> <li>2. Abstract (1 page)</li> </ol> <p><i>(Rationale of the study, preliminary results, detailed description of the translational value of the research and the expected impact on the NHS)</i></p> <ol style="list-style-type: none"> <li>3. introduction</li> <li>4. background and rationale</li> <li>5. experimental design (organized in tasks)</li> <li>6. further details on the overall methods that will be used in this project</li> <li>7. work carried out and preliminary results</li> <li>8. expected results and relevant corresponding milestones</li> <li>9. References and relevant publications by the research group, already available</li> </ol> <p><i>(max 8 pages)</i></p>
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PERSONNEL INVOLVED IN THE RESEARCH				
Name and date of birth	Role on Project	Fellowship required	Effort on project (%)	Present position

<p>DESCRIPTION OF THE WORK FOR EVERY UNIT OF PERSONNEL</p> <ol style="list-style-type: none"> <li>1.....</li> <li>2.....</li> <li>3.....etc.....</li> </ol> <p>Budget Form /year</p> <ol style="list-style-type: none"> <li>1. research costs</li> </ol>
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2. Instruments 3. Indirect costs <b>4. Sub-total.....</b> 5. Overheads 6. Fellowships <b>7. Total.....</b>
Justifications Itemized research costs.....
<b>EXISTING/PENDING SUPPORT.....</b>
<b>SUGGESTED REVIEWERS (MAX 3)</b> ..... ..... .....
<b>BIOETHICAL REQUIREMENT</b>  1. Human experimentation .....(YES/NOT) – please provide clearance from the competent ethical committee as <u>addendum A</u>  2. Animal experimentation .....(YES/NOT) – please include a statement as <u>addendum B</u> specifying which regulations the proposed research meets
<b>Declaration</b> I shall confirm to the Declaration of Helsinki in its latest version. I shall also apply the Bioethics Convention of the Council of Europe. In implementing the proposed research, I shall adhere most strictly to all existing ethical and safety provisions applicable. Before start of the research, I shall obtain clearance from the competent ethical committee in case of involvement of human subjects in the research and /or in case of other ethical implications. I shall conform with all regulations protecting the animals used for research purpose.  Date: ..... Name of PI.....signature

**Principal investigator's signature** .....

**Authorized Administrative Official's signature**.....

Date

Si autorizza al trattamento dei dati ai sensi del d.lgs. 196/2003